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### INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Complete if Known **Application Number** 09/547,506 Filing Date April 12, 2000 First Named Inventor Christian, S.T. Group Art Unit **Examiner Name** Attorney Docket Number IMI-001

OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the Cite Examiner item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), Initials Nο publisher, city and/or country where published. Alexander, N., Yoneda, S., Vlachakis, N.D. and R.F. Maronde. 1984. Role of conjugation and red blood cells for inactivation of ciculating catecholamines. Am. J. Physiol. 247 (1): R203-R207 Alvarado, F. and R.K. Crane. 1960. Phlorizin as a competitive inhibitor of the active transport of sugars by hamster small intestine, in vitro. Biochim. Biophys. Acta 56: Arita, H. and J. Kawanami. 1980. Studies on uptake of phenyl glycosides as inhibitors of 3 D-glucose uptake by Rhesus monkey kidney cells. J. Biochem. 88: 1399-1406. Barnett, J.E.G., Holman, G.D. and K.A. Munday. 1973. Structural requirements for binding to the sugar transport system of the human erythrocyte. Biochem. J. 131: 211-221. Barnett, A., McQuade, R.D. and C. Tedford. 1992. Highlights of D1 dopamine receptor antagonist research. Neurochem. Int. 20 (Suppl.): 119S-122S. Bencsics, A., Sershen, H., Baranyi, M., Hashim, A., Lajtha, A. and E.S. Vizi. 1997. 6 Dopamine, aswell as, norepinephrine, is a link between noradrenergic nerve terminals and splenocytes. Brain Res. 761 (2): 236-243. Berger, J.G., Chang, W.K., Clader, J.W., Hou, D., Chipkin, R.E. and A.T. McPhail. 1989. Synthesis and receptor affinities of some conformationally restricted analogues of the dopamine D1 selective ligand (5R)-8-chloro-2,3,4,5-tetrahydro-3-methyl-5-phenyl- 1H-3-benzazepin-7-ol. J. Med. Brewster, W.K., Nichols, D.E., Riggs, R.M., Mottola, D.M., Lovenberg, T.W., Lewis, M.H. and R.B. Mailman. 1990. trans-10,11-dihydroxy-5,6,7,8,12b-hexahydrobenzo[a]phen -anthridine: A highly potent selective dopamine D1 full agonist. J. Med. Chem. 33: 1756-1764. Bodor, N., Roller, R.G. and S.J. Selk. 1978. Elimination of a quaternary pyridinium salt delivered as its dihydropyridine. J. Pharm. Sci, 67 (5): 685 Bodor, 1976. "Novel Approaches for the Design of Membrane Transport Properties of Drugs". In: "Design of Biopharmaceutical Properties Through Prodrugs and Analogs", Ed. E.B. Roche et al. APhA Academy of Pharmaceutical Sciences, Washington, D.C., pp. 98-135 Bodor, N., Farag, H.H. and M.E. Brewster. 1981. Site-specific, sustained release of drugs to the brain. Science 214: 1370-1372

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Examiner Initials	Cite No.	OTHER PRIOR ART — NON PATENT LITERATURE DOCUMENTS  Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T²
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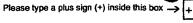
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Filing Date	April 12, 2000						
First Named Inventor	Christian, S.T.						
Group Art Unit							
Examiner Name							
Attorney Docket Number	IMI-001						

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Application Number 09/547,506

Filing Date April 12, 2000

First Named Inventor Christian, S.T.

Group Art Unit

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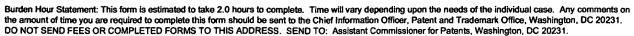
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09/547,506 **Application Number Filing Date** April 12, 2000 First Named Inventor Christian, S.T. Group Art Unit **Examiner Name** 

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Attorney Docket Number IMI-001

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